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(57) Abstract: This invention concerns the compounds of formula (I), prodrugs, <I>N</I>-oxides, addition salts, quaternary amines, metal complexes or stereochemically isomeric forms thereof wherein -a1-a2-a3-a4- is a radical of formula -CH=CH-CH=CH-, -N=CH-CH=CH-, -CH=N-CH=CH-, -CH=CH-N=CH-, -CH=CH-CH=N- wherein each hydrogen atom may optionally be substituted; Q is a radical of formula (b-1), (b-2), (b-3), (b-4), (b-5), (b-6), (b-7), (b-8), wherein Alk is C₁₋₆alkanediyl; Y1 is a bivalent radical of formula-NR2- or -CH(NR2R4)-; X1 is NR4, S, S(=O), S(=O)2, O, CH2, C(=O), CH(=CH2), CH(OH), CH(CH₃), CH(OCH₃), CH(SCH₃), CH(NR^{5a}R^{5b}), CH₂-NR⁴ or NR⁴-CH₂; X² is a direct bond, CH₂, C(=O), NR⁴, C₁₋₄alkyl-NR⁴, NR⁴-C₁₋₄alkyl; t is 2 to 5; u is 1 to 5; v is 2 or 3; and whereby each hydrogen in Alk and in (b-3), (b-4), (b-5), (b-6), (b-7) and (b-8), may optionally be replaced by R3; provided that when R3 is hydroxy or C1-calkyloxy, then R3 can not replace a hydrogen atom in the α position relative to a nitrogen atom; G is substituted $C_{1:10}$ alkanediyl wherein the substituent is attached via an oxygen atom; R^1 is an optionally substituted monocyclic heterocycle or aryl; R2 is hydrogen, formyl, C1.6alkylcarbonyl, Hetcarbonyl, pyrrolidinyl, piperidinyl, homopiperidinyl, C3-7cycloalkyl or C1-10alkyl substituted with N(R6)2 and optionally with another substituent; R3 is hydrogen, hydroxy, C_{1-6} alkyl, C_{1-6} alkyloxy, aryl C_{1-6} alkyl or aryl C_{1-6} alkyloxy; R^4 is hydrogen, C_{1-6} alkyl or aryl C_{1-6} alkyl; R^{5a} , R5b, R5c and R5d are hydrogen or C1-6alkyl; or R5a and R5b, or R5c and R5d taken together form a bivalent radical of formula C(CH₂),- wherein s is 4 or 5; R⁶ is hydrogen, C₁₋₄alkyl, formyl, hydroxyC₁₋₅alkyl, C₁₋₅alkylcarbonyl or C₁₋₅alkyloxycarbonyl; aryl is optionally substituted phenyl; Het is pyridyl, pyrimidinyl, pyrazinyl, pyridazinyl; as respiratory syncytial virus replication inhibitors; their preparation, compositions containing them and their use as a medicine.



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